California Medical Device Recall Information

Recall Name
Baxter Recalls VASCU-Guard Peripheral Vascular Patch Due to Potential for Incorrect Patch Orientation

Recall Date	Product Description	Recalling Firm	Recall Reason
05/02/15	VASCU-Guard Peripheral Vascular Patch	**Baxter International, Inc.** Deerfield, IL	*Baxter received customer complaints of difficulty in distinguishing the smooth from rough surface of the VASCU-GUARD patch.* *Incorrect orientation of the patch with the rough side toward the bloodstream may increase the risk of vessel thrombosis and/or embolism.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	• Vascu-Guard TS, 1x6cm **Product code 1504026** • Vascu-Guard TS, 0.8x8cm **Product code 1504028** • Vascu-Guard TS, 1x10cm **Product code 1504030** • Vascu-Guard TS, 2x9cm **Product code 1504032**	**CA**, nationwide	Manufacturing dates: January 19, 2015 to May 1, 2015 Distribution dates: March 16, 2015 to May 1, 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm449825.htm